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AMENDMENTS TO THE CLAIMS:

Listing of Claims:

This listing of claims will replace all prior versions of the claims and listing of the claims in the application:

(Currently Amended) A method for treating a subject for a disease treatable by modulation of RNA (DTMR) associated with splicing of nuclear RNA, comprising: administering to said subject an effective amount of a tetracycline compound of formula (I):

wherein

 R^2 , R^2 , R^4 , and R^4 are each independently hydrogen[[,]] or alkyl-alkenyl, alkynyl, alkylthio, alkylsulfinyl, alkylsulfonyl, alkylamino, arylalkyl, aryl, heterocyclic, heteroaromatic or a prodrug moiety;

 $R^3, R^{10}, R^{11} \ and \ R^{12} \ are \ each \ hydrogen, \\ alkyl, \ alkenyl, \ alkynyl, \ substituted \\ earbonyl, \ or \ a \ pro-drug \ moiety;$

R4 is NR4'R4", alkyl, alkenyl, alkynyl, hydroxyl, halogen, or hydrogen;

 R^{S} is hydroxyl, hydrogen, thiol, alkanoyl, aroyl, alkaroyl, aryl, heteroaromatic, alkyl, alkenyl, alkynyl, alkoxy, alkylthio, alkylsulfinyl, alkylsulfonyl, alkylamino, arylalkyl, alkyl carbonyloxy, or aryl-carbonyloxy;

 $R^6 \ and \ R^6' \ arc \ cach \ independently-hydrogen, absent, hydroxyl, halogen, thiel, alkyl, alkenyl, alkynyl, aryl, alkoxy, alkylthio, alkylsulfinyl, alkylsulfonyl, alkylamino, or an arylalkyl;$

 $R^7 \ is \ \textit{hydrogen, hydroxyl, halogen, thiol, nitro, alkyl, substituted_alkenyl, substituted_alkenyl, substituted_phenyl, substituted or unsubstituted furanyl, alkexy, a$

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alkylthio, alkylsulfinyl, alkylsulfonyl, arylalkyl, amino, arylalkenyl, arylalkynyl, acyl, or aminoalkyl, heterocyclic, thionitroso, or -(CH₂)_{0.3}NR^{7e}C(-W')WR^{7e};

R⁸ is hydrogen, hydroxyl, halogen, thiol, nitro, alkyl, alkenyl, alkynyl, aryl, alkoxy, alkylthio, alkylsulfinyl, alkylsulfonyl, alkylamino, amino, arylalkenyl, arylalkynyl, acyl, aminoalkyl, heterocyclic, thionitroso, or—(CH₂)_{0.3}NR[®]C(=E')ER^{®a};

 $R^{9} \ is \ hydrogen, hydroxyl, \ halogen, thiol, nitro, alkyl, alkenyl, alkynyl, aryl, alkoxy, alkylthio, alkylsulfinyl, alkylsulfonyl, arylalkyl, amino, arylalkenyl, arylalkynyl, aeyl, aminoalkyl, heterocyclic, thionitroso, or <math>-(CH_{3})_{0.3}NR^{9}(C=Z')ZR^{9};$ and

 $R^{2a}, R^{2b}, R^{2c}, R^{2d}, R^{2c}, R^{2f}, R^{8b}, R^{8b}, R^{8c}, R^{8c}, R^{8c}, R^{8c}, R^{9c}, R^{9$

 $R^{13} : is hydrogen, hydroxy, alkyl, alkenyl, alkynyl, alkoxy, alkylthio, aryl, alkylsulfinyl, alkylsulfonyl, alkylamino, or an arylalkyl; alkylsulfonyl, al$

Y² and Y are each independently hydrogen, halogen, hydroxyl, cyano; sulfhydryl, amino, alkyl, alkenyl, alkynyl, alkoxy, alkylthio, alkylsulfinyl, alkylsulfonyl, alkylamino, or an arylalkyl;

Z' is O₂ S₂ or NR ^{of}₂ or a pharmaceutically acceptable salt, ester or enantiomer thereof₂ with the proviso that said tetracycline compound is not tetracycline;

such that said DTMR associated with splicing of nuclear RNA is treated, wherein said DTMR associated with splicing of nuclear RNA is spinal muscular atrophy, and further

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wherein said effective amount is effective to modulate splicing of said subject's nuclear RNA

2.-36. (Canceled)

The method of claim 1, wherein R²[[,]] and R²'₇ 37. (Previously Presented) R8-R10-R11-and R12 are each hydrogen. X is CR6R62-and R4 is NR42R42, wherein and R4 and R4" are each methyl.

38. (Canceled)

- The method of claim 38 claim 37, wherein R7 is 39. (Currently Amended) substituted or unsubstituted arylfuranyl.
- The method of elaim 39 claim 37, wherein R7 is 40. (Currently Amended) substituted-or-unsubstituted phenyl.

41. (Canceled)

42. The method of elaim-41 claim 40, wherein said (Currently Amended) substituted phenyl is substituted with one or more substituents and further wherein said substituents are each independently alkyl, alkenyl, alkynyl, halogen, hydroxyl, alkylcarbonyloxy, arylcarbonyloxy, alkoxycarbonyloxy, aryloxycarbonyloxy, carboxylate, alkylcarbonyl, arylcarbonyl, alkoxycarbonyl, aminocarbonyl, alkylaminocarbonyl, dialkylaminocarbonyl, alkylthiocarbonyl, alkoxyl, phosphate, phosphonato, phosphinato, cyano, amino, acylamino, amidino, imino, sulfhydryl, alkylthio, arylthio, thiocarboxylate, sulfates, alkylsulfinyl, sulfonato, sulfamoyl, sulfonamido, nitro, trifluoromethyl, cyano, azido, heterocyclyl, alkylaryl, aryl or heterocyclic moiety.

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43. (Currently Amended) The method of elaim-38claim 37, wherein R⁷ is substituted or unsubstituted alkenyl.

- 44. (Currently Amended) The method of claim 37, wherein \mathbb{R}^7 is substituted alkynylor-unsubstituted heteroaryl and \mathbb{R}^9 is alkyl.
- 45. (Currently Amended) The method of elaim-36 claim 37, wherein R^7 is dialkylaminoacyl.
- 46. (Currently Amended) The method of elaim 45 claim 37, wherein $\mathbb{R}^9 \underline{\mathbb{R}^7}$ is alkylaminoaminoalkyl.

47.-56. (Canceled)

57. (Previously Presented) The method of claim 1, wherein said tetracycline compound is:

(Canceled)

 (Previously Presented) The method of claim 1, wherein said modulation of splicing increases splicing of RNA. Applicants: Levy, et al. Serial No.: 10/692,764 Filing Date: October 24, 2003 Page -6-

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- (Previously Presented) The method of claim 1, wherein said modulation of splicing decreases splicing of RNA.
 - 61. (Canceled)
- 62. (Previously Presented) The method of claim 1, wherein said subject is a mammal.
- 63. (Previously Presented) The method of claim 62, wherein said mammal is a human.
- 64. (Previously Presented) The method of claim 1, wherein said modulation of splicing is activation of cryptic splice sites, silencing of consensus splice sites, silencing of exonic or intronic splicing enhancers (ESEs or ISEs), silencing of exonic or inronic splicing silencers (ESEs or ISSs), alteration of the binding or a component of the splicing machinery to the RNA, or the affecting of intermolecular interactions between components of the splicing machinery.
- 65. (Previously Presented) The method of claim 1, wherein said tetracycline compound is:

or a pharmaceutically acceptable salt thereof.

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66. (New) A method for treating a subject for a DTMR associated with splicing of nuclear RNA, comprising: administering to said subject an effective amount of a tetracycline compound; wherein said tetracycline compound is a tetracycline compound selected from the group consisting of:

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and pharmaceutically acceptable salts, esters and enantiomers thereof; such that said DTMR associated with splicing of nuclear RNA is treated, wherein said DTMR associated with splicing of nuclear RNA is spinal muscular atrophy, and further wherein said effective amount is effective to modulate splicing of said subject's nuclear RNA.